INTER-OFFICE MEMORANDUM

TO R. Hiatt, M.D.
AT Division of Research

DATE November 22, 1989

FROM Glenda R. Marlow

Glenda R. Marlow

AT KFRI Administration
3505 Broadway
Oakland

COPIES TO E. Van Brunt, M.D.

SUBJECT Institutional Review and Approval of Research Study Entitled "Pregnancy Outcome Study"

On November 21, 1989, the Institutional Review Board (IRB) for the protection of human subjects reviewed the above-subject research study. The study was approved for continuation as amended subject to revision of the consent form and receipt and approval of the questionnaire acceptable to Staff Counsel. These will be sent to you as soon as they have been reviewed and approved. Enclosed is a statement entitled "Information About Rights of Medical Research Participants." A copy of this statement must be given to each prospective participant in order to satisfy statutory requirements.

All continuing projects and activities must be reviewed and reapproved at least annually by the Board. Board approval of any project is for a maximum period of one year. It is the responsibility of the investigator to resubmit the project for periodic review as determined by the Board. You will be reminded of the scheduled follow-up review dates for your project.

If this proposal is used in conjunction with any other human experimentation or if it is modified in any way, it must be reapproved for these special circumstances. In addition, the Board must be promptly notified of unanticipated problems involving risks to human subjects or any complications which may occur during any experimental procedure.

If you have any questions, please call me at ext. 3236.

GRM/mlm
Enclosure

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Source: https://www.industrydocuments.ucsf.edu/docs/zpmm0000